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=> file reg COST IN U.S. DOLLARS	SINCE FILE	TOTAL SESSION
FULL ESTIMATED COST	58.29	250.05
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)		TOTAL
CA SUBSCRIBER PRICE	-7.65	-7.65

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STRUCTURE FILE UPDATES: 15 MAR 2010 HIGHEST RN 1210111-73-1 DICTIONARY FILE UPDATES: 15 MAR 2010 HIGHEST RN 1210111-73-1

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TSCA INFORMATION NOW CURRENT THROUGH January 8, 2010.

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REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/support/stngen/stndoc/properties.html

=> Uploading c:\program files\stnexp\queries\10540993 3.17.10

15 STRUCTURE UPLOADED

=> s 15

SAMPLE SEARCH INITIATED 11:51:07 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 467 TO ITERATE

100.0% PROCESSED 467 ITERATIONS SEARCH TIME: 00.00.01 14 ANSWERS

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 8044 TO 10636

McIntosh

PROJECTED ANSWERS: 56 TO 504

14 SEA SSS SAM L5

14 ANSWERS REGISTRY COPYRIGHT 2010 ACS on STN Adenosine, N=[(4-chloro-2-nitrophenyl)methyl]- (9CI) C17 H17 Cl N6 O6 1.6

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):3

14 ANSWERS REGISTRY COPYRIGHT 2010 ACS on STN Adenosine, N-[(2,4-dihydroxyphenyl)methyl]- (9CI) C17 H19 N5 O6 IN

Absolute stereochemistry.

MF

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

14 ANSWERS REGISTRY COPYRIGHT 2010 ACS on STN Adenosine, N-[(4-hydroxy-2,5-dimethoxyphenyl)methyl)- (9CI) H23 N5 07 IN

MF

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L6 14 ANSWERS REGISTRY COPYRIGHT 2010 ACS on STN Adenosine, N-[(2,3,6-trifluorophenyl)methyl]-MF C17 H16 F3 N5 O4

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

- => d 15 L5 HAS NO ANSWERS
- L5 STF
- * STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY AVAILABLE VIA OFFLINE PRINT * Structure attributes must be viewed using STN Express query preparation.
- -> d his

(FILE 'HOME' ENTERED AT 10:51:09 ON 17 MAR 2010)
FILE 'REGISTRY' ENTERED AT 10:51:21 ON 17 MAR 2010

```
STRUCTURE UPLOADED
             14 S L1
             249 S L1 FULL
     FILE 'CAPLUS' ENTERED AT 10:52:11 ON 17 MAR 2010
             48 S L3
     FILE 'REGISTRY' ENTERED AT 11:50:44 ON 17 MAR 2010
                STRUCTURE UPLOADED
1.6
-> s 15 full
FULL SEARCH INITIATED 11:55:57 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED -
                                     8800 TO ITERATE
                                                                     242 ANSWERS
100.0% PROCESSED
                      8800 ITERATIONS
SEARCH TIME: 00.00.01
             242 SEA SSS FUL L5
-> file caplus
COST IN U.S. DOLLARS
                                                      SINCE FILE
                                                                      TOTAL
                                                           ENTRY
                                                                     SESSION
FULL ESTIMATED COST
                                                          195.95
                                                                      446.00
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)
                                                     SINCE FILE
                                                                       TOTAL
                                                          ENTRY
                                                                     SESSION
CA SUBSCRIBER PRICE
                                                             0.00
                                                                        -7.65
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FILE COVERS 1907 - 17 Mar 2010 VOL 152 ISS 12
FILE LAST UPDATED: 16 Mar 2010 (20100316/ED)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Dec 2009
USPTO MANUAL OF CLASSFICATIONS THESAURUS ISSUE DATE: Dec 2009
CAplus now includes complete International Patent Classification (IPC)
reclassification data for the first quarter of 2010.
CAS Information Use Policies apply and are available at:
http://www.cas.org/legal/infopolicy.html
This file contains CAS Registry Numbers for easy and accurate
substance identification.
           192 L7
=> d bib abs hitstr 180-192
     ANSWER 180 OF 192 CAPLUS COPYRIGHT 2010 ACS on STN
AN
     1974:121282 CAPLUS
    80:121282
DM
OREF 80:19535a,19538a
     2',3',5'-Tri-O-acyl-N6-benzyladenosines
     Kampe, Wolfgang; Fauland, Erich; Thiel, Max; Roesch, Egon; Dietmann, Karl
IN
Dh
    Boehringer Mannheim G.m.b.H.
    Ger. Offen., 12 pp.
```

CODEN: GWXXBX

Patent

T.A German

FAN	.CNT 1 PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 2238923	A1.	19740214	DE 1972-2238923	19720808
	CA 1003411	A1	19770111	CA 1973-177826	19730731
	GB 1384518	A	19750219	GB 1973-36489	19730801
	AU 7358857	A	19750206	AU 1973-58857	19730802
	CH 579587	A5	19760915	CH 1973-11307	19730803
	FR 2195434	A1	19740308	FR 1973-28648	19730806
	ZA 7305331	A	19740828	ZA 1973-5331	19730806
	NL 7310870	A	19740212	NL 1973-10870	19730807
	AT 7306918	A	19750115	AT 1973-6918	19730807
	AT 325784	В	19751110		
	JP 49045095	A	19740427	JP 1973-89161	19730808

PRAI DE 1972-2238923 A GI For diagram(s), see printed CA Issue.

For dragram(s), see printed CA issue. Eight acyladenosines I (R = Ac, Bz, or nicotinoyl, Rnl = 2-Me, 2,5-Me2, 2,4,5-Me3, 2,5-Me0Cl, or 2,5-MeSCl) were prepared in 45-85% yield by acylation of I (R = B) with Ae2O, BCL, or nicotinoyl azide. The acyl derivs. had longer lasting effects on blood vessels and circulation than AB the starting compds. I (R = H). 23707-33-7 34349-31-0 34349-38-7

52622-05-6

RL: RCT (Reactant); RACT (Reactant or reagent)

(acylation of) 23707-33-7 CAPLUS Adenosine, N-[(2-methylphenyl)methyl]- (CA INDEX NAME)

Absolute stereochemistry.

34349-31-0 CAPLUS Adenosine, N-[(2,5-dimethylphenyl)methyl]- (9CI) (CA INDEX NAME)

RN

34349-38-7 CAPLUS Adenosine, N-[[5-chloro-2-(methylthio)phenyl]methyl]- (9CI) (CA INDEX CN

Absolute stereochemistry.

52622-05-6 CAPLUS Adenosine, N-[(2,4,5-trimethylphenyl)methyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

OSC.G 2 THERE ARE 2 CAPLUS RECORDS THAT CITE THIS RECORD (2 CITINGS) ANSWER 181 OF 192 CAPLUS COPYRIGHT 2010 ACS on STN

```
1974:93268 CAPLUS
AN
      Cytokinins in Populus x robusta. Light effects on endogenous levels
ΑU
      Hewett, E. W.; Wareing, P. F.
      Dep. Bot. Microbiol., Univ. Coll. Wales, Aberystwyth, UK
Planta (1973), 114(2), 119-29
GODEN: PLANAR; 1583N: 0032-0935
SO
       English
AB
      Cytokinin levels in both attached and detached mature leaves of poplar (P.
       robusta) increased transiently after short periods of exposure to red
      light. The degree and rapidity of response seems dependent on the physiol. condition of the leaves. The cytokinin,
      6\text{-}(2\text{-hydroxybenzyl}) aminopurine riboside, specifically increased after red light treatment. Diurnal changes of leaf cytokinins occurred, with a
```

pronounced peak of activity being present at daybreak. 50868-58-1 RL: BIOL (Biological study)

(of poplar, red light effect on)

50868-58-1 CAPLUS

Adenosine, N-((2-hydroxyphenyl)methyl)- (CA INDEX NAME)

Absolute stereochemistry.

OSC.G 8 THERE ARE 8 CAPLUS RECORDS THAT CITE THIS RECORD (8 CITINGS)

```
L8
    ANSWER 182 OF 192 CAPLUS COPYRIGHT 2010 ACS on STN
```

1973:534312 CAPLUS AN

DN 79:134312

OREF 79:21771a,21774a

New cytokinin from Populus robusta

AU

Horgan, R.; Hewett, E. W.; Purse, J. G.; Wareing, P. F. Dep. Bot. Microbiol., Univ. Coll. Wales, Aberystwyth, UK Tetrahedron Letters (1973), (30), 2827-8

CODEN: TELEAY; ISSN: 0040-4039

Journal

LA English

For diagram(s), see printed CA Issue,

A new cytokinin was isolated from the leaves of P. robusta and shown to be AB $6-[(o-hydroxybenzyl)amino]-9-\beta-D-ribofuranosylpurine (I).$ 50868-58-1

RL: BIOL (Biological study)

(in Populus robusta)

50868-58-1 CAPLUS RN

Adenosine, N-[(2-hydroxyphenyl)methyl]- (CA INDEX NAME)

ANSWER 183 OF 192 CAPLUS COPYRIGHT 2010 ACS on STN AN 1973:413413 CAPLUS

DN 79:13413

OREF 79:2119a,2122a

Inhibitors of nucleoside and nucleotide metabolism

ĀŪ Henderson, J. F.; Paterson, A. R. P.; Caldwell, I. C.; Paul, B.; Chan, M. C.; Lau, K. F.

CS Cancer Res. Unit, Univ. Alberta, Edmonton, AB, Can.

Cancer Chemotherapy Reports, Part 2 (1973), 3(1), 71-85 CODEN: CCSUBJ; ISSN: 0069-0120 so

Journal LA English

A total of 164 purine and pyrimidine derivs. and analogs were screened for inhibition of nucleoside and nucleotide metab in 4 test systems. Among a

number of potent inhibitors identified, No-(3-methyl-2-butenyl)-adenosine [7724-76-7] and 4-(dimethylamino)-7-B-D-ribofuranosyl-7H-pyrrolo[2,3-dipyrimidine [I] [20371-00-0] inhibited de novo purine biosynthesis in

incubated Ehrlich ascites tumor cells,

ac(-amino-9-y1)-a'-(hydroxymethyl)diglycolaldehyde-bis(phenylhydrazone) (II) [40297-82-7] inhibited adenine phosphoribosyltransferase [9027-80-9] from Ehrlich ascites tumor cells, 4-amino-5-iodo-7-β-D-ribofuranosyl-7H-pyrrolo[2,3-d]pyrimidine
[24386-93-4] inhibited adenine kinase [9027-72-9] activity in tumor cell

exts., and 2-amino-6-[(p-fluorobenzyl)thio]-9-β-D-ribofuranosyl-9Hpurine (III) [40297-53-8] and N6-(p-nitrobenzyl)-adenosine [40297-54-9] inhibited nucleoside transport (inosine synthesis) in

incubated human erythrocytes.

40297-54-9 RL: BIOL (Biological study)

(inosine formation by erythrocytes in response to)

RN 40297-54-9 CAPLUS CN

Adenosine, N-[(4-nitrophenyl)methyl]- (9CI) (CA INDEX NAME)

- ANSWER 184 OF 192 CAPLUS COPYRIGHT 2010 ACS on STN
- 1973:124846 CAPLUS AN DN
- OREF 78:20071a,20074a
- N-Benzyladenosine derivatives
- TN Kampe, Wolfgang; Fauland, Erich; Thiel, Max; Juhran, Wolfgang; Stork,
- PA Boehringer Mannheim G.m.b.H.
- Ger. Offen., 20 pp. CODEN: GWXXBX SO
- Patent
- LA German

FAN.	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 2136624	A	19730208	DE 1971-2136624	19710722
	GB 1340643	A	19731212	GB 1972-33537	19720618
	US 3845035	A	19741029	US 1972-271098	19720712
	ZA 7204891	A	19730530	ZA 1972-4891	19720717
	CH 569035	A5	19751114	CH 1975-10617	19720719
	CH 570420	A5	19751215	CH 1972-10795	19720719
	NL 7210023	A	19730124	NL 1972-10023	19720720
	CA 979891	A1	19751216	CA 1972-147625	19720720
	SU 539532	A.3	19761215	SU 1972-1812966	19720720
	FR 2146493	A1	19730302	FR 1972-26450	19720721
	AT 317446	В	19740826	AT 1972-6288	19720721
	AT 790673	A	19750415	AT 1973-7906	19720721
PRA:	I DE 1971-2136624	A	19710722		

- For diagram(s), see printed CA Issue.
 - Thirty-three title compds. (I; X = NHCH2C6H5-nRn; R: = Cl, OH NH2 or Br; AB Rn = e.g. 2-OH, 3,2-HOMe, 2,5 HOCl, 2,4-HOCl) were prepared by reaction of I (X - Cl) containing free or acetyl group-protected OH-groups with H2NCH2C6H5-nRn or from the adenosine derivative and ClCH2C6H5nRn. I had circulatory and antilipemic effects.
 - 40896-26-2P 40297-54-9P 40896-32-0P 40896-39-7P 40896-40-0P 40896-41-1P
 - 40896-43-3P 40896-45-5P 40896-50-2P
 - 40896-52-4P 40958-96-1P 40958-97-2P
 - RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of)
 - 40297-54-9 CAPLUS RN
- Adenosine, N-[(4-nitrophenyl)methyl]- (9CI) (CA INDEX NAME) CN

40896-26-2 CAPLUS

Adenosine, N-[[3-(hydroxymethyl)-2-methylphenyl]methyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 40896-32-0 CAPLUS

Adenosine, N-[[5-(hydroxymethyl)-2-methylphenyl]methyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

40896-39-7 CAPLUS Adenosine, N-[[2-(hydroxymethyl)-5-methylphenyl]methyl]- (9CI) (CA INDEX CN

NAME)

Absolute stereochemistry.

RN

40896-40-0 CAPLUS Adenosine, N-[(2-nitrophenyl)methyl]- (CA INDEX NAME)

Absolute stereochemistry.

RN 40896-41-1 CAPLUS

Adenosine, N-[(5-methyl-2-nitrophenyl)methyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

40896-43-3 CAPLUS RN

CN Adenosine, N-[(2-methoxy-5-nitrophenyl)methyl]- (9CI) (CA INDEX NAME)

- 40896-45-5 CAPLUS Adenosine, N-[(2-methyl-3-nitrophenyl)methyl]- (9CI) (CA INDEX NAME) CN

Absolute stereochemistry.

- 40896-50-2 CAPLUS Benzonitrile, 3-[[(9- β -D-ribofuranosyl-9H-purin-6-yl)amino]methyl]-(CA INEXNAME)

- 40896-52-4 CAPLUS RN
- Adenosine, N-[(4-cyanophenyl)methyl]- (CA INDEX NAME)

40958-96-1 CAPLUS Adenosine, N-[(3-nitrophenyl)methyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

40958-97-2 CAPLUS Adenosine, N-[(3-cyano-2-methylphenyl)methyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

THERE ARE 3 CAPLUS RECORDS THAT CITE THIS RECORD (3 CITINGS) OSC.G 3

- ANSWER 185 OF 192 CAPLUS COPYRIGHT 2010 ACS on STN
- AN 1972:502140 CAPLUS 77:102140
- DN
- N-[[(Hydrazinocarbonyl)phenyl]alkyl]adenosines
- Jahn, Werner; Kampe, Wolfgang; Fauland, Erich; Juhran, Wolfgang; Stork,
- PA Boehringer Mannheim G.m.b.H.
- Ger. Offen., 14 pp. CODEN: GWXXBX so
 - Patent
- German FAN.CNT 1

1.151.0	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 2060189	A	19720615	DE 1970-2060189	19701208
	US 3787391	A	19740122	US 1971-201174	19711122
	NL 7116564	A	19720612	NL 1971-16564	19711202
	GB 1313459	A	19730411	GB 1971-56025	19711202
	SU 444368	A3	19740925	SU 1971-1721738	19711202
	AU 7136492	A	19730607	AU 1971-36492	19711203
	CH 567045	A5	19750930	CH 1971-17640	19711203
	CH 568330	A5	19751031	CH 1975-8284	19711203
	CH 568331	A5	19751031	CH 1975-8285	19711203
	ZA 7108177	A	19720927	ZA 1971-8177	19711207
	HU 163227	В	19730728	HU 1971-B01335	19711207
	AT 312172	В	19731227	AT 1971-10533	19711207
	AT 318821	В	19741125	AT 1972-9168	19711207
	AT 318822	В	19741125	AT 1972-9169	19711207
	CA 960656	A1	19750107	CA 1971-129590	19711207
	FR 2117935	A5	19720728	FR 1971-43996	19711208
	FR 2117935	B1	19750314		
	SU 515454	A3	19760525	SU 1973-1959114	19730824
	SU 576955	A3	19771015	SU 1973-1959113	19730824
PRA	I DE 1970-2060189	A	19701208		

For diagram(s), see printed CA Issue.

- Fourteen title compds. (I, 2-, 3-, 4-, or 5-CONHNHR1; Q CH2, CH2CH2, CH2CH2O; R = H, 2-Me, 3-Cl; Ri = H, p-clC6H4CO, p-MeoC6H4CO, p-MeoC6H4CO, p-MeoC6H4CO, useful as blood-circulation-active and serum-lipids-lowering agents, were prepared by reaction of
 - tri-O-acetyladenosine with R(RINHNHCO)C6H3QBr or of adenosine N-[R(Eto2C)C6H3Q] derivative with N2H4.H2O. 38790-46-4P 38790-49-7P 38790-52-2P
- (preparation of) RN
- RL: SPN (Synthetic preparation); PREP (Preparation) Benzoic acid, 4-methyl-3-[[(9-β-D-ribofuranosyl-9H-purin-6
 - yl)amino]methyl]-, hydrazide (CA INDEX NAME)

38790-49-7 CAPLUS

Benzoic acid, 2-methyl-3-[[(9-\beta-D-ribofuranosyl-9H-purin-6yl)amino]methyl]-, hydrazide (CA INDEX NAME)

Absolute stereochemistry.

RN 38790-52-2 CAPLUS

Benzoic acid, 3-methyl-[[(9-\$-D-ribofuranosyl-9H-purin-6yl)amino]methyl]-, hydrazide (9CI) (CA INDEX NAME)

Absolute stereochemistry.

THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD (1 CITINGS)

ANSWER 186 OF 192 CAPLUS COPYRIGHT 2010 ACS on STN 1972:502139 CAPLUS 77:102139 AN

DN

OREF 77:16847a,16850a

N-(Acylbenzyl- and -phenethyl)adenosines Kampe, Wolfgang; Fauland, Erich; Stork, Harald; Juhran, Wolfgang; IN Dietmann, Karl

PA Boehringer Mannheim G.m.b.H.

SO Ger. Offen., 20 pp. CODEN: GWXXBX

Patent

German

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI	DE	2059922	A	19720615	DE 1970-2059922	19701205
	US	3817981	A	19740618	US 1971-199727	19711117
	SU	469253	A3	19750430	SU 1971-1723201	19711130
	SU	506294	A3	19760305	SU 1971-1913745	19711130
	NL	7116563	A	19720607	NL 1971-16563	19711202
	GB	1313290	A	19730411	GB 1971~56024	19711202
	CH	567044	A5	19750930	CH 1971-17633	19711202
	CH	573445	A5	19760315	CH 1975-8318	19711202
	FR	2116517	A5	19720713	FR 1971-43419	19711203
	FR	2116517	B1	19750801		
	ZA	7108104	A	19720927	ZA 1971-8104	19711203
	AU	7136493	A	19730607	AU 1971-36493	19711203
	HU	163670	В	19731027	HU 1971-B01334	19711203
	AT	314094	В	19740325	AT 1971-10436	19711203
	CA	960655	A1	19750107	CA 1971-129319	19711203
	AT	323335	В	19750710	AT 1971-323335	19711203
PRAI	DE I	1970-2059922	A	19701205		

DE 170-203922 A 170-203922 For diagram (s), see printed CM Issue.
Forty-five title compds. (I, Y = X,2-R(R1)C6H39CH2)nNH; n = 1,2; R = 3 - or AΒ 4-carboxy, -alkoxycarbonyl, -carbamoyl, -allylcarbamoyl; R1 = H, Me; R2 = H, Cl, OH) (II), useful as hypolipemic agents with effects on circulation, were prepared by reaction of the corresponding I (Y - Cl) (III) with X, $2-R(R) | C6H2 \rangle nNH2$ and subsequent saponification or amidation. Thus, refluxing III (R2-H) and 3-EtO2C-C6H4CH2CH2NH2.HC1 in EtOH in the presence of ENN for 3 hr gave 65% II (n = 2, R = 3-EO2C, RI = R2 = H), which was heated in EtOH at 120° for 15 hr with NH3 to give 64% II (n = 2, R = 3-EO2C, RI = R2 = H), which was heated in EtOH at 120° for 15 hr with NH3 to give 64% II (n = 2, R = 3-EO2C), RI = R2 = 5h), 38623-36-26 38623-36-27 38623-39-5p

38823-69-7P 38823-66-4P 38823-81-3P 38823-82-4P 38823-90-4P

RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of)

38823-50-6 CAPLUS

Benzoic acid, 2-methyl-3-[[(9-β-D-ribofuranosyl-9H-purin-6-yl)amino]methyl]-, ethyl ester (CA INDEX NAME)

Absolute stereochemistry.

38823-56-2 CAPLUS

Benzoic acid, 4-methyl-3-[[(9-β-D-ribofuranosyl-9H-purin-6yl)amino]methyl]-, ethyl ester (CA INDEX NAME)

- 38823-59-5 CAPLUS
- Benzoic acid, 3-methyl-4-[[(9-β-D-ribofuranosyl-9H-purin-6-yl)amino]methyl]-, ethyl ester (CA INDEX NAME)

- 38823-66-4 CAPLUS
 Benzoic acid, 2-methyl-3-[[(9-B-D-ribofuranosyl-9H-purin-6-yl)amino]methyl]- (CA INDEX NAME)

RN 38823-69-7 CAPLUS

N Benzoic acid, 4-methyl-3-[[(9-B-D-ribofuranosyl-9H-purin-6-yl)amino]methyl]- (CA INDEX NAME)

Absolute stereochemistry.

RN 38823-72-2 CAPLUS CN Benzoic acid, 3-met

N Benzoic acid, 3-methyl-4-[[(9-β-D-ribofuranosyl-9H-purin-6-yl)amino]methyl]- (CA INDEX NAME)

Absolute stereochemistry.

RN 38823-79-9 CAPLUS CN Adenosine, N-[[5-(aminocarbony1)-2-methylpheny1]methyl]- (9CI) (CA INDEX NAME)

38823-81-3 CAPLUS

Adenosine, N-[[3-(aminocarbonyl)-2-methylphenyl]methyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

38823-82-4 CAPLUS Adenosine, N-[[4-(aminocarbonyl)-2-methylphenyl]methyl]- (9CI) (CA INDEX NAME)

38823-90-4 CAPLUS

Adenosine, N-[[2-methyl-3-[(methylamino)carbonyl]phenyl]methyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

OSC.G 4 THERE ARE 4 CAPLUS RECORDS THAT CITE THIS RECORD (5 CITINGS)

1.8 ANSWER 187 OF 192 CAPLUS COPYRIGHT 2010 ACS on STN

AN 1972:483708 CAPLUS

DN

OREF 77:13769a,13772a

- Clinical-pharmacological studies with a new orally active adenosine derivative
 - Schaumann, E.; Kutscha, W. Sonadum, 2., Actsona, M. Iniv. Heidelberg, Mannheim, Fed. Rep. Ger. Arzneimittel-Forshung (1972), 22(4), 783-90 CODEN: ARXND; ISSN: 0004-4175.
- - Journal
- T.A

ΑU

CS SO

German Metrifudil [N6-(o-methylbenzyl)adenosine] (I) [23707-33-7] was AB tested in humans. Administration of 0.03 mg/kg i.v. and of 0.35 mg/kg orally increased the heart rate and cardiac output. Neither impairment of atrioventricular conduction nor other alterations of the electrocardiogram was observed Uneasiness and other side effects were caused by i.v. and oral administration of 0.1 and 0.47-0.53 mg I/kg, resp. The limit of tolerability was reached earlier if the speed of i.v. infusion exceeded 16 $\mu g/kg/min$. No critical changes in circulatory parameters were found. I.v. injection of I caused no inflammation or alteration of the veins. The concentration of serum fatty acids was lowered only by i.v. administration of

- I. A 50% absorption of I was estimated by comparing the increase of the heart rate after i.v. and oral administration.
- RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BTOL (Biological study); USES (Uses)
- (pharmacol. of) 23707-33-7 CAPLUS
- Adenosine, N-[(2-methylphenyl)methyl]- (CA INDEX NAME)

Absolute stereochemistry.

- OSC.G 5 THERE ARE 5 CAPLUS RECORDS THAT CITE THIS RECORD (5 CITINGS)
- ANSWER 188 OF 192 CAPLUS COPYRIGHT 2010 ACS on STN
- 1972:154069 CAPLUS AN
- 76:154069 DM
- OREF 76:25121a,25124a
 - Novel synthesis of N6-substituted adenosines and their coronary dilator activities
- Shimizu, Bunji; Kaneko, Masakatsu; Saito, Akio; Nishino, Hiroshi; Mizuno,
- Hiroshi; Nakayama, Koichi; Ohshima, Takeshi; Koike, Hiroyuki Sankyo Res. Lab., Tokyo, Japan Sankyo Kenkyusho Nenpo (1971), 23, 117-23
- SO
- CODEN: SKKNAJ: ISSN: 0080-6064
- Journal Japanese
- NG-Substituted adenosine derivs. (PhCH2, PhCH2CH2, naphthylmethyl, Me2CHCH2, o-MeC6H4-CH2, m-MeC6H4CH2, p-MeC6H4CH2, furfurylmethyl) in addition
 - to N6-benzyl-9-(β-D-arabinofuranosyl)adenine, and N6-benzyl-9-(β-D-glucopyranosyl) adenine were synthesized directly from adenosine by exchange amination reactions of the corresponding purine or pyrimidine bases. The mechanism of formation of these nucleosides and
- their coronary-dilating activities were described. 23707-33-7P 35940-03-5P 35940-04-6P
- RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of, as blood vessel dialators)
- 23707-33-7 CAPLUS Adenosine, N-[(2-methylphenyl)methyl]- (CA INDEX NAME)

RN

35940-03-5 CAPLUS Adenosine, N-[(3-methylphenyl)methyl]- (CA INDEX NAME) CN

Absolute stereochemistry.

RN

35940-04-6 CAPLUS Adenosine, N-[(4-methylphenyl)methyl]- (CA INDEX NAME)

Absolute stereochemistry.

ANSWER 189 OF 192 CAPLUS COPYRIGHT 2010 ACS on STN 1971:541121 CAPLUS 75:141121 L8 AN

DN

- Coronary dilating N6-benzyladenosines Kampe, Wolfgang; Fauland, Erich; Thiel, Max; Dietmann, Karl; Juhran,
- Wolfgang
- PA Boehringer Mannheim G.m.b.H.
- Ger. Offen., 10 pp. CODEN: GWXXBX
- Patent

J.A.	German	
DAM	CMT 1	

FAN.	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 2007273	A	19710826	DE 1970-2007273	19700218
	SU 399134	A3	19730927	SU 1971-1616102	19710129
	US 3781273	A	19731225	US 1971-112424	19710203
	NL 7102026	A	19710820	NL 1971-2026	19710216
	DK 123357	В	19720612	DK 1971-694	19710216
	HU 162739	В	19730428	HU 1971-B01274	19710216
	CH 549596	A	19740531	CH 1971-2208	19710216
	CH 549600	A	19740531	CH 1974-2849	19710216
	CA 953714	A1	19740827	CA 1971-105563	19710216
	ZA 7101030	A	19711124	ZA 1971-1030	19710217
	FR 2081524	A5	19711203	FR 1971-5318	19710217
	FR 2081524	B1	19740927		
	AT 306251	В	19730410	AT 1971-1378	19710217
	AT 313483	В	19740225	AT 1972-1233	19710217
	JP 51016440	В	19760524	JP 1971-7691	19710218
	GB 1279946	A	19720628	GB 1971-1279946	19710419
PRAI	DE 1970-2007273	A	19700218		

For diagram(s), see printed CA Issue.

The title compds. (I, where R - Me, MeS, or MeO, R1 - 5-Me, 5-Cl, 5-MeO, 5-iso-Pr, 5-F, 5-tert-Bu, 3-Me, or 3-Cl) were prepared wither by amination AB

of the 6-chloro derivative or by N1-substitution of adenosine followed by alkaline

rearrangement. Thus, 9-(2,3,5-ti-0-acetyl-B-D-ribofunanosyl)-6-chloropurine, 2-may 2

5-Me). Similarly prepared were 11 other I. 34349-31-0P 34349-32-1P 34349-33-2P 34349-34-3P 34349-35-4P 34349-37-6P

34349-38-7P 34349-39-8P 34349-41-2P 34422-72-5P

RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of)

34349-31-0 CAPLUS

Adenosine, N-[(2,5-dimethylphenyl)methyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

34349-32-1 CAPLUS

Adenosine, N-[5-methyl-2-(methylthio)benzyl]- (8CI) (CA INDEX NAME)

10/540,993

34349-33-2 CAPLUS RN

Adenosine, N-(5-chloro-2-methylbenzyl)- (8CI) (CA INDEX NAME) Absolute stereochemistry.

34349-34-3 CAPLUS Adenosine, N-(5-methoxy-2-methylbenzyl)- (8CI) (CA INDEX NAME) Absolute stereochemistry.

34349-35-4 CAPLUS Adenosine, N-(2-methoxy-5-methylbenzyl)- (8CI) (CA INDEX NAME) Absolute stereochemistry.

34349-37-6 CAPLUS Absolute stereochemistry.

Adenosine, N-[(5-fluoro-2-methylphenyl)methyl]- (9CI) (CA INDEX NAME)

RN

34349-38-7 CAPLUS Adenosine, N-[[5-chloro-2-(methylthio)phenyl]methyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

34349-39-8 CAPLUS

Adenosine, N-(5-tert-buty1-2-methylbenzy1)- (8CI) (CA INDEX NAME)

RN 34349-40-1 CAPLUS CN Adenosine, N-[(2,3-dimethylphenyl)methyl]- (9CI) (CA INDEX NAME) Absolute stereochemistry.

RN 34349-41-2 CAPLUS
CN Adenosine, N-[(3-chloro-2-methylphenyl)methyl)- (9CI) (CA INDEX NAME)
Absolute stereochemistry.

RN 34422-72-5 CAPLUS CN Adenosine, N-(5-isopropyl-2-methylbenzyl)- (8CI) (CA INDEX NAME) Absolute stereochemistry.

osc.g 4 THERE ARE 4 CAPLUS RECORDS THAT CITE THIS RECORD (4 CITINGS)

ANSWER 190 OF 192 CAPLUS COPYRIGHT 2010 ACS on STN an

1971:433660 CAPLUS DN

OREF 75:5316h,5317a

Pharmacological effects on coronary reactive hyperemia in conscious dogs

Junran, W.; Voss, E. M.; Dietmann, K.; Schaumann, W. Pharmakol. Lab., Boehring Mannheim G.m.b.H., Mannheim, Fed. Rep. Ger. AU

CS

so Naunyn-Schmiedebergs Archiv fuer Pharmakologie (1971), 269(1), 32-47 CODEN: NNAPBA; ISSN: 0340-5249

Journal.

LA English

For diagram(s), see printed CA Issue. In conscious dogs, threshold doses of dipyridamole (I) and lidoflazine (II), which potentiate the dilation of coronary vessels by adenosine, increased reactive hyperemia in response to arterial occlusion lasting >30 sec, whereas threshold doses of coronary dilators, such as N6-(o-methylbenzyl) adenosine (III) and carbochromen (IV), which do not potentiate adenosine, did enhance reactive hyperemia for any duration of occlusion. Theophylline decreased the duration of reactive hyperemia, but not the excess flow. Procaine-HCl infused into the coronary artery caused a dose-dependent reduction of the reactive hyperemia. Apparently, appreciable amts. of adenosine were liberated only during complete anoxia for >30 sec. Under physiol. conditions coronary resistance was probably regulated by a nervous mechanism and not by adenosine liberation.

RL: BIOL (Biological study) (hyperemia response to)

RN 23707-33-7 CAPLUS

Adenosine, N-[(2-methylphenyl)methyl]- (CA INDEX NAME)

Absolute stereochemistry.

OSC.G 2 THERE ARE 2 CAPLUS RECORDS THAT CITE THIS RECORD (2 CITINGS)

- ANSWER 191 OF 192 CAPLUS COPYRIGHT 2010 ACS on STN AN 1971:86054 CAPLUS
- 74 86054
- OREF 74:13963a,13966a
- Inhibition of induced thrombocyte aggregation by adenosine and adenosine derivatives. II. Correlation between inhibition of the aggregation and
- ΑU Dietmann, Karl; Birkenheier, H.; Schaumann, Wolfgang
- Med. Forsch., Firma Boehringer Mannheim G.m.b.H., Mannheim-Waldhof, Fed. Rep. Ger.
- Arzneimittel-Forschung (1970), 20(11), 1749-51 CODEN: ARZNAD; ISSN: 0004-4172
- German
- For diagram(s), see printed CA Issue.
- AB The ability of adenosine (I) and 20 adenosine derivs. to produce vasodilation in rabbits was correlated with their ability to antagonize ADP-induced thrombocyte aggregation in vitro. The N6-phenylalkyl substituted derivs., N6-(cis, trans-2-phenylcyclo-pentyl)adenosine and N6-(trans-dl-2-phenylcyclopentyl)adenosine (II), were more active than the aliphatic substituted derivs., 2-chloro-N6-propyl-, 2-chloro-N6-allyl-,
 - and 2-chloro-N6-sec-butyladenosines, as well as the N6-benzyl derivs., 2-chloro-N6-benzyladenosine, 2-amino-N6-(2-chlorobenzyl)adenosine, N6-(o-xylyl)adenosine, N6-(o-trifluoromethylbenzyl)adenosine, and N6-(3.5-dimethoxybenzyl)adenosine. The most active derivative, II, was half as active as adenosine.
- RL: BIOL (Biological study)
- (blood platelet aggregation and vasodilation by) 23707-33-7 CAPLUS
- Adenosine, N-[(2-methylphenyl)methyl]- (CA INDEX NAME)

- ANSWER 192 OF 192 CAPLUS COPYRIGHT 2010 ACS on STN AN 1969:115505 CAPLUS
- DN 70:115505
- OREF 70:21591a,21594a
- N6-Aralkyl adenosine derivatives IN
- Thiel, Max; Stach, Kurt; Jahn, Werner; Schaumann, Wolfgang; Dietmann, Karl Boehringer, C. F., und Soehne G.m.b.H. S. African, 15 pp. PA
- 80
- CODEN: SFXXAB Patent
- A English

FAN.	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	ZA 6707414		19680502		
	DE 1670171			DE	
	FR 1550512			FR	
	GB 1145789			GB	

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US 3506643
                                      19700414
                                                                               19671018
PRAI DE
                                      19661209
      DE
                                      19670711
      MARPAT 70:115505
      For diagram(s), see printed CA Issue.
      The title compds. (1), where halogen, alkyl, alkoxy, F3C or alkylthio, or
     two substituents may be H or a methylenedioxy, are prepared from the corresponding D-ribosides and benzylamines, or from the corresponding
      N'-substituted adenosine derivs. Thus, 8.2 q.
      tri-O-acetyl-6-chloro-9-β-D-ribosyl-9-H-purine and 7.2 g.
      2-ClC6H4CH2NH2 in 120 cc. iso-PrOH were refluxed 2 hrs., worked up and the
      residue dissolved in 100 cc. MeOH, 10 cc. N NaOH solution added and the mixture
      refluxed 1 hr. to yield 4 q. I (R = 2-Cl), m. 182-3°. The
      following I were similarly prepared (R and m.p. given): 3,4-Cl2,
     182-3°; 4-MeO, 146-7°; 3,4 (MeO) 2, 135-6°; 4-Cl, 174-5°; 3,4,5-(MeO) 3, 118-19°; 2,6-Cl2, 207-9°; 4-Cl, 174-5°;
      3-C1, 168-9°; 2-MeO, 147-8°; 2-Me, 157-8°; 3,5-(MeO)2, 191-2°; 2-MeS, 127-8°; 2-F3C, 160-1°; and
      2',3'-0-isopropylideneadeno-sine in 200 cc. MeCN, 10 g. p-BrC6H4Br was
      added and the mixture refluxed 24 hrs. with stirring. The precipitate which formed
      was filtered off, dissolved in 150 cc. MeOH and an equal volume 2N NaOH
      solution was added. The mixture was heated on a steam bath 20 min., extracted with
      CHC13, evaporated, and the residue dissolved in 200 cc. HC02N. Water was
      added until the mixture became cloudy. The mixture was left standing 1 day at
      ambient temperature, after which it was evaporated in vacuo, and the residue made weakly alkaline with an aqueous solution of concentrated NH3 to yield 5.8 q. I (R = 4-Br),
     m. 168-9°. I exhibit an effect on blood vessels and circulation. 23661-00-9P 23707-33-7P
     RL: SPN (Synthetic preparation); PREP (Preparation)
     (preparation of)
23661-00-9 CAPLUS
     Adenosine, N-[o-(methylthio)benzyl]- (8CI) (CA INDEX NAME)
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Absolute stereochemistry.

RN 23707-33-7 CAPLUS

CN Adenosine, N-[(2-methylphenyl)methyl]- (CA INDEX NAME)

OSC.G 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD (1 CITINGS)